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(54) Title: VARIANTS OF HUMAN PAPILLOMA VIRUS ANTIGENS			
(57) Abstract Variants of human papilloma virus (HPV) E6 and E7 proteins able to elicit a humoral and/or cellular immune response against HPV in a host animal but not being cell-transforming in the host animal are disclosed, and are useful in treatment or prevention of diseases or conditions involving HPV.			

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VARIANTS OF HUMAN PAPILLOMA VIRUS ANTIGENS

FIELD OF THE INVENTION

5

This invention relates generally to variants of human papilloma virus (HPV) antigens, and in particular it relates to non-transforming variants of HPV antigens which are suitable for use in vaccines. The invention also extends to vaccine compositions which include these variants of HPV antigens as active immunogens, as well as to
10 methods of use of these variants to elicit an immune response against HPV.

BACKGROUND OF THE INVENTION

Papillomaviruses are small DNA viruses that infect a variety of animal species.
15 Some are associated with the development of malignancies in their natural hosts. Over 60 types of human papillomavirus (HPV) have been identified. These infect humans at a variety of body locations and are responsible for common skin warts, laryngeal papillomas, genital warts and other wart-like lesions. Genital HPV infections are particularly common and a number of HPV types, but most frequently types 6, 11, 16 and
20 18, infect the genital tract in both men and women. In women, HPVs infect various portions of the genital tract including the cervix.

Genital HPVs are a significant clinical problem. HPV infection of the ano-genital region is now regarded as the most common form of viral sexually-transmitted disease
25 (STD). The viruses cause genital infections which become manifest in one of three ways:

- i clinical infection, where gross genital warts are visible;
- ii subclinical infection, where viral lesions are not obvious but are detectable using special viewing techniques; and
- iii latency, where the only sign of infection is the presence of HPV DNA.

30

Subclinical infections are common. It is estimated that 2 to 4% of Papanicolaou (Pap.) smears show evidence of HPV. Latent infections are even more frequent and the majority of adults harbour one or more types of genital HPV.

5 Carcinoma of the uterine cervix (CaCx) is a common cancer in women. Two forms of cervical cancer are recognised; squamous cell carcinoma (SCC) is by far the most frequent representing about 90% of observed cases; adenocarcinoma, a cancer of the secretion cells, accounts for about 10%. Cancer of the cervix develops through pre-cancerous intermediate stages to invasive forms (the carcinoma) which can become life
10 threatening. The pre-cancerous stages of increasing severity are known as cervical intraepithelial neoplasia (CIN) grades 1 to 3. Over a 20 year period about 40% of the untreated CIN3 patients develop invasive cancer, the increasingly serious forms of which are known as stage I to IV. Invasive cancer frequently leads to death.

15 Cervical cancer in both its pre-cancerous and invasive stages is one of the few cancers for which a highly reliable and relatively cheap screening method is available. The Papanicolaou (Pap.) smear involves cytological examination of cervical scrapes to test for the presence of abnormal cervical cells which are indicative of pre- or invasive cancer. Detection of abnormalities leads to further investigation and treatment if
20 necessary.

To be effective at reducing the number of cervical cancers and resultant deaths, Pap. smear screening is undertaken on a mass scale and ideally includes all women of sexually-active age. Detection and subsequent treatment of CIN has a very high success
25 rate in the prevention of invasive cancer, while early detection of the latter can have a marked effect on mortality.

Most developed countries have highly developed Pap. smear screening programs which have resulted in a 30% drop in age-specific mortality due to CaCx between 1960
30 and 1980. However, apart from the Scandinavian countries, few developed countries screen more than 50 to 60% of women, allowing CaCx and resultant deaths to remain a significant problem. In the developing world the situation is even worse as few organised

screening programs exist, resulting in 400,000 new cases of invasive cancer annually in these countries.

As outlined earlier, a variety of types of HPV cause genital infections in humans, although four types (6, 11, 16, 18) predominate. Evidence collected over the past 15 years strongly suggests that several of the HPVs are associated with the development of cervical cancer. Indeed many researchers have concluded that specific HPV types are the essential aetiologic factor responsible for the development of many of the cancers.

10 Infection with HPV-16 and HPV-18 has been associated with the development of cancer of the cervix. It has been postulated that HPV acts as an initiator in cervical carcinogenesis and that malignant transformation depends on interaction with other factors. Infections with HPV-6 and HPV-11 have been associated with the development of genital warts. The incidence of HPV infection appears to be increasing as shown by
15 a large increase recently in patient visits related to genital HPV infections in both males and females and the presence of HPV in Pap. smears of some women under 30 years of age.

The nature of HPV-16 in particular and papilloma viruses in general has been well
20 studied recently. HPV-16 contains a 7904 bp double-stranded DNA genome (Siedorf, K. *et al.*, *Virology* (1985) 145:181-185). The capsid is 50 nm and contains 72 capsomers (Klug, A., *J. Mol. Biol.* (1965) 11:403-423). U.S. Patent 4,777,239 discloses a series of 17 synthetic peptides which are said to be capable of raising antibodies to HPV-16 and thus may be useful for diagnostic purposes. In addition, European Patent 0 412 762
25 discloses polypeptides which are antagonists of the biochemical interaction of the HPV E7 protein and the retinoblastoma gene (RBG) protein, and which are said to be useful in the treatment of genital warts and cervical cancer.

The DNAs of several papilloma viruses have been sequenced, including several
30 HPV types, bovine papillomavirus (BPV) and cottontail rabbit papillomavirus (CRPV). All of these display similar patterns of nucleotide sequence with respect to open reading frames. The open reading frames can be functionally divided into early regions (E) and

late regions (L); the E regions are postulated to encode proteins needed for replication and transformation; and the L regions to encode the viral capsid proteins (Danos, O., *et al.*, *J. Invest. Derm.* (1984) 83:7s-11s).

5 Two HPV encoded proteins, E6 and E7, are thought to be involved in the pathogenesis of HPV-induced abnormal cell proliferation (reviewed in Stoppler *et al.*, *Intervirology*, (1994) 37:168-179). The amino acid sequences of the HPV-16 E6 and E7 proteins as deduced from the nucleic acid sequence are shown in Siedorf *et al.*, *Virology*, (1985) 145:181-185.

10

The HPV genes encoding the E6 and E7 proteins are invariably expressed in tissue or tumor cells obtained from cervical cancers associated with HPV infection. In addition, the HPV E6 and E7 genes derived from the HPV-16 strain are capable of inducing epithelial cell transformation in cell culture without the presence of other HPV genes.

15 These observations indicate that at least part of the stimulation of cell proliferation caused by HPV infection is due to the E6 and E7 viral proteins.

The HPV E6 and E7 proteins are believed to be effective immunological targets for tumour regression. As described above, however, the E6 and E7 genes are known to

20 "transform" cells possibly by the action of their protein products in interfering with cellular proteins involved in the control cell growth. Accordingly, if even minute traces of DNA encoding the E6 and E7 proteins were to be present in a vaccine preparation, this could cause that vaccine preparation to initiate irreversible transformation events in the cells of a recipient of the vaccine preparation. It is an object of the present invention to

25 provide non-transforming variants of the HPV E6 and E7 proteins which are able to induce in a host animal (particularly a human) a range of humoral and cellular immune responses, and which are therefore suitable for use in the production of vaccines for the prevention, prophylaxis, therapy and treatment of HPV-induced diseases or other conditions which would benefit from inhibition of HPV infection.

30

In the work leading to the present invention, it has been recognised that there are four ways to induce immune responses to E6 and/or E7 proteins:

- (i) use whole proteins (this introduces the possibility that contaminating DNA may be associated with the proteins);
- (ii) use point mutants (this can lead to reversion to native protein, which requires multiple mutations to avoid; in addition, any point mutation leads to loss of potentially vital epitopes);
- (iii) use specific peptides (this requires a very large number of peptides, the identification of which is very complex, to make a vaccine of broad utility); and
- (iv) use variants such as fusions and combinations of deletion mutants (this method has none of the above limitations).

In addition to the cell transforming properties of the E7 protein itself, fusions of this protein with β -galactosidase have also been shown to be cell-transforming (Fujikawa *et al.*, *Virology*, 204, 789-793, 1994). Accordingly, it could not be predicted that fusions of E6 and/or E7 moieties, either full length or non-full length, would not also be cell-transforming.

SUMMARY OF THE INVENTION

In one aspect the present invention provides as an isolated protein, a variant of the HPV E6 or E7 protein which variant is able to elicit a humoral and/or cellular immune response against HPV in a host animal but which is not cell-transforming in the host animal.

In another aspect, the present invention provides a method for eliciting an immune response against HPV in a host animal, which method comprises administering to the host animal an effective amount of a variant of the HPV E6 or E7 protein which variant is able to elicit a humoral and/or cellular immune response against HPV in a host animal but which is not cell-transforming in the host animal.

In yet another aspect, the present invention provides a vaccine composition for use in eliciting an immune response against HPV in a host animal, which comprises a variant

of the HPV E6 or E7 protein which variant is able to elicit a humoral and/or immune response against HPV in a host animal but which is not cell-transforming in the host animal, and optionally an adjuvant, together with a pharmaceutically acceptable carrier and/or diluent.

5

The invention also extends to the use of a variant of the HPV E6 or E7 protein which variant is able to elicit a humoral and/or cellular immune response against HPV in a host animal but which is not cell-transforming in the host animal, and optionally an adjuvant, in eliciting an immune response against HPV in a host animal.

10

Throughout this specification, unless the context requires otherwise, the word "comprise", or variations such as "comprises" or "comprising", will be understood to imply the inclusion of a stated integer or group of integers but not the exclusion of any other integer or group of integers.

15

DETAILED DESCRIPTION OF THE INVENTION

As used herein, references to the variant of this invention as being "not cell-transforming in the host animal" mean that the cell-transforming property of the "parent" or wild-type HPV E6 or E7 protein has been reduced, and preferably effectively eliminated, in the variant. In particular, these references indicate that this cell-transforming property has been significantly reduced in comparison with wild-type E6 or E7 protein in appropriate test systems.

20

It will be appreciated that where the non-transforming variant HPV E6 or E7 protein of this invention is produced by expression of an appropriate encoding recombinant DNA molecule, the nature of that encoding DNA, unlike the wild-type E6 or E7 genes, would not have the potential to initiate irreversible transformation events in the cells of the host animal.

25

30

The variant HPV E6 or E7 proteins of the present invention include, but are not limited to, deletion mutants of the wild-type E6 or E7 proteins in the form of non-full

length fragments of the wild-type proteins, as well as fusion proteins in which E6 and/or E7 moieties are fused, optionally with a linkage of from 1 to 50, preferably a short linkage of from 1 to 20, and more preferably from 1 to 5, amino acid residues between the E6 and/or E7 moieties. The E6 and/or E7 moieties in such a fusion protein may
5 comprise the full wild-type E6 or E7 proteins, or alternatively they may comprise non-full length fragments of the wild-type proteins. The fusion proteins may also comprise other moieties fused or otherwise coupled thereto, for example moieties to assist in purification of the fusion protein (for example, a glutathione-S-transferase or GST moiety or hexa-His moiety) or to enhance the immunogenicity of the fusion protein (for example an adjuvant
10 such as diphtheria or cholera toxin or a non-toxic derivative thereof such as the holotoxoid or B sub-unit of cholera toxin).

The term "non-full length fragment" is used herein to describe polypeptides which may for example comprise deletion mutants of the E6 or E7 proteins corresponding to at
15 least 50%, more preferably 60-70%, and even 80-90% of the full-length E6 or E7 protein sequence. By way of example only, the fragments may be deletion mutants corresponding to the N-terminal or C-terminal two-thirds of the E6 or E7 proteins.

Suitable non-full length fragments, and fusion proteins which comprise the E6
20 and/or E7 proteins or non-full length fragments thereof, as described above may be readily produced by techniques which are well known in the art and which are described by way of example below. It will be appreciated by persons skilled in this art that variant HPV E6 or E7 proteins as described above including fusion proteins which comprise various combinations of the E6 and/or E7 moieties may be readily produced using these
25 known techniques, and then tested using routine methods to establish whether the resultant fusion protein or other variant protein meets the criteria of the present invention, that is whether it is able to elicit a humoral and/or cellular immune response in a host animal but is not cell-transforming in the host animal.

30 Preferably, the host animal is a human, however the host animal may also be a non-human mammal.

The present invention is particularly, but not exclusively, directed to variants of the E6 or E7 proteins of the HPV-16 and HPV-18 genotypes, however it will be appreciated that the invention extends to variants of the corresponding proteins in other HPV genotypes, particularly the HPV-6 and HPV-11 genotypes which are causative agents of condylomata acuminata, and other genotypes which have oncogenic potential of a type similar to HPV-16 and HPV-18.

Previous work in this area has shown that vaccination of rats with live viral vectors expressing HPV E6 or E7 proteins leads to rejection of transplanted E7-bearing tumour cells (Meneguzzi *et al.*, *Virology*, 181:62-69, 1991), while vaccination of cattle with an adjuvanted HPV E7 vaccine leads to accelerated rejection of tumours induced by bovine papillomavirus (Campo, *Cancer Cells*, 3:421-426, 1991).

The variant HPV E6 or E7 proteins of the present invention are provided as isolated proteins, that is they are substantially free of other HPV proteins, and find particular utility for the treatment of genital warts, cervical cancer or other conditions caused by HPV in man. The variant proteins can be included in pharmaceutical compositions for the treatment or prevention of diseases involving HPV as well as the other conditions discussed above.

The variant HPV E6 or E7 proteins of the invention may be used to raise antibodies and/or induce cellular immune responses, either in subjects for which protection against infection by HPV is desired, i.e. as prophylactic vaccines, or to heighten the immune response to an HPV infection already present, i.e. as therapeutic vaccines. They also can be injected into production species to obtain antisera. In lieu of the polyclonal antisera obtained in the production species, monoclonal antibodies may be produced using the standard methods or by more recent modifications thereof by immortalising spleen or other antibody-producing cells for injection into animals to obtain antibody-producing clones. The polyclonal or monoclonal antibodies obtained, corrected if necessary for species variations, can also be used as therapeutic agents.

Direct administration of the variant proteins to a host can confer either protective immunity against HPV or, if the subject is already infected, a boost to the subject's own immune response to more effectively combat the progress of the HPV induced disease.

5 The magnitude of the prophylactic or therapeutic dose of a variant HPV E6 or E7 protein of this invention will, of course, vary with the group of patients (age, sex, etc.), the nature or the severity of the condition to be treated and with the particular variant protein and its route of administration. In general, the weekly dose range for use lies within the range of from about 0.1 to about 5 μ g per kg body weight of a mammal.

10

Any suitable route of administration may be employed for providing a mammal, especially a human, with an effective dosage of a variant protein of this invention. For example, oral, rectal, vaginal, topical, parenteral, ocular, nasal, sublingual, buccal, intravenous and the like may be employed. Dosage forms include tablets, troches,
15 dispersions, suspensions, solutions, capsules, creams, ointments, suppositories, aerosols and the like. Said dosage forms also include injected or implanted slow releasing devices specifically designed for this purpose or other forms of implants modified to additionally act in this fashion.

20 If the variant proteins are to be administered as vaccines, they are formulated according to conventional methods for such administration to the subject to be protected. If the antibodies are to be used for therapeutic purposes, it is generally desirable to confer species characteristics upon them compatible with the subject to be treated. Accordingly, it is often desirable to prepare these antibodies in monoclonal form since fusion with
25 suitable partners is capable of conferring the desired characteristics on the secreted monoclonals.

The variant proteins may be delivered in accordance with this invention in ISCOMS™ (immune stimulating complexes), liposomes or encapsulated in compounds
30 such as acrylates or poly(DL-lactide-co-glycoside) to form microspheres. The variant proteins may also be incorporated into oily emulsions and delivered orally.

Other adjuvants, as well as conventional pharmaceutically acceptable carriers, excipients, buffers or diluents, may also be included in the vaccine compositions of this invention. Generally, a vaccine composition in accordance with the present invention will comprise an immunologically effective amount of the variant HPV E6 or E7 protein, and
5 optionally an adjuvant, in conjunction with one or more conventional pharmaceutically acceptable carriers and/or diluents. An extensive though not exhaustive list of adjuvants can be found in Coulter and Cox, "Advances in Adjuvant Technology and Application", in *Animal Parasite Control Utilizing Biotechnology*, Chapter 4, Ed. Young, W.K., CRC Press, 1992. As used herein "pharmaceutically acceptable carriers and/or diluents" include
10 any and all solvents, dispersion media, aqueous solutions, coatings, antibacterial and antifungal agents, isotonic and absorption delaying agents and the like. The use of such media and agents for pharmaceutical active substances is well known in the art and is described by way of example in *Remington's Pharmaceutical Sciences*, 18th Edition, Mack Publishing Company, Pennsylvania, U.S.A.

15

In practical use, a variant protein of this invention can be combined as the active ingredient in intimate admixture with a pharmaceutical carrier according to conventional pharmaceutical compounding techniques. The carrier may take a wide variety of forms depending on the form of preparation desired for administration, e.g. oral or parenteral
20 (including intravenous and intra-arterial). In preparing the compositions for oral dosage form, any of the usual pharmaceutical media may be employed, such as, for example, water glycols, oils, alcohols, flavouring agents, preservatives, colouring agents and the like in the case of oral liquid preparations, such as, for example, suspensions, elixirs and solutions; or carriers such as starches, sugars, microcrystalline cellulose, diluents,
25 granulating agents, lubricants, binders, disintegrating agents and the like in the case of oral solid preparations such as, for example, powders, capsules and tablets. Because of their ease of administration, tablets and capsules represent the most advantageous oral dosage unit form, in which case solid pharmaceutical carriers are obviously employed. If desired, tablets may be sugar-coated or enteric-coated by standard techniques.

30

In addition to the common dosage forms set out above, the variant proteins of this invention may also be administered by controlled release means and/or delivery devices,

including by way of example, the controlled release preparations disclosed in International Patent Specification No. PCT/AU93/00677 (Publication No. WO 94/15636).

Pharmaceutical compositions of the present invention suitable for oral or parenteral administration may be presented as discrete units such as capsules, cachets or tablets each containing a predetermined amount of the active ingredient, as a powder or granules or as a solution or a suspension in an aqueous liquid, a non-aqueous liquid, an oil-in-water emulsion or a water-in-oil liquid emulsion. Such compositions may be prepared by any of the methods of pharmacy but all methods include the step of bringing into association the active ingredient with the carrier which constitutes one or more necessary ingredients. In general, the compositions are prepared by uniformly and intimately admixing the active ingredient with liquid carriers or finely divided solid carriers or both, and then, if necessary, shaping the product into the desired presentation.

Further features of the present invention are more fully described in the following Example(s). It is to be understood, however, that this detailed description is included solely for the purposes of exemplifying the present invention, and should not be understood in any way as a restriction on the broad description of the invention as set out above.

EXAMPLE 1

Cloning and expression of GST E6/E7 fusion protein.

A molecule consisting of HPV-16 E6 and E7 sequences as an "in-frame" fusion was created as follows. A clone of HPV-16 DNA containing both E6 and E7 genomic sequences served as the template for separate PCR amplification of E6 and E7 using oligonucleotides:

- (a) (5')CGCTCGAGAGATCTCATATGCACCAAAAGAGAACTGC(3') and
(b) (5')CGCCCGGGCAGCTGGGTTTCTCTACGTG(3') for E6;
and

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- (c) (5')CGCCCGGGATGCATGGAGATACACCTACATTGCATG(3') and
(d) (5')CGGTCGACGGATCCTGGTTTCTGAGAACAGATGGG(3') for E7.

A *Sma*I recognition site at the 3' end of E6 and the 5' end of E7 facilitated the fusion and introduced two additional amino acids (proline and glycine) between E6 and E7. Additional restriction enzyme recognition sites at the 5' and 3' boundaries of the fusion molecule (introduced in the oligonucleotides) aided in subsequent cloning procedures.

10 The fused E6/E7 sequence was cloned as a *Bgl*II-*Bam*HI fragment into vector pDS56 (Stuber *et al.*, *EMBO J.*, (1984) 3:3143-3148) which provided an in-frame 3' hexa-his(hh) sequence. From this, E6/E7hh was removed as a *Eco*RI/*Hind*III fragment and subcloned into pGEM 7 + 3, which was created by inserting the *Bam*HI/*Hind*III portion of the pGEM3-Zf(+)(Promega) polylinker into the *Bam*HI/*Hind*III site of the multiple cloning site of the pGEM7-Zf(+)(Promega) vector. E6/E7hh was then removed from pGEM7+3 as a *Eco*RI/*Sal*I fragment and inserted into the multiple cloning site of pGEX-4T-1 (Pharmacia) to produce pGEX-4T-1 E6/E7hh. This plasmid was used to transform a variety of *E. coli* strains including TOPP2 (Stratagene) and BL21 (Amrad/Pharmacia). Both types of transformed cells produced a significant amount of fusion protein following IPTG induction (Fig. 2). The fusion protein (GST E6/E7hh represented schematically in Fig. 1a) was in the expected size range of around 60kDa. The identity of the protein was confirmed by Western blots probed with two monoclonal antibodies directed against E7 (LHIL.16E7.8F and LHIL.16E7.6D, Tindle *et al. Journal of General Virology*, (1990) 71:1347-1354) (Fig. 3).

25

EXAMPLE 2

Cloning and expression of E6/E7 fusion protein

30 In order to express E6/E7hh as protein lacking GST, a termination codon was introduced into pGEX-4T-1 E6/E7hh at a unique *Ba*II site 3' to, and in-frame with, the GST translation initiation codon using the phosphorylated linker TGCTCTAGAGCA.

After transforming *E. coli* strain BL21 with this new plasmid ([GST] E6/E7hh) a significant amount of protein (E6/E7hh, represented schematically in Fig.1b) was produced following IPTG induction at a size of approximately 33kD which corresponds to the size expected of a E6/E7hh fusion protein (Fig. 2). Identity of this protein was confirmed by Western blot using the same monoclonal antibodies as in Example 1 (Fig. 3).

EXAMPLE 3

10 Cloning and expression of deleted (non-full length) forms of E6 and E7

(i) Construction of Δ E6C/ Δ E7N

Full length E6/E7 in pGEM3(Promega) served as a template for PCR amplification of deleted forms of E6/E7 using oligonucleotides 5'GCGCGAATTCTATTAAGGAGCCCGGGATGGGGAATCCATATGCTGT-AT3' and 5'CGCGAGATCTCCGAAGCGTAGAGTCACACTTG3'.

The resulting truncation of E6/E7 lacking sequences (189bp) at the N terminal of E6 and C terminal of E7 (96bp) was subcloned into pGEX-4T-1 containing a termination codon in the GST sequence to produce [GST] Δ E6C/ Δ E7Nhh. This plasmid was used to transform *E. coli* strain BL21. Transformed cells expressed a significant amount of fusion protein (Δ E6C/ Δ E7Nhh, represented schematically in Fig.1c) following IPTG induction (Fig. 4a) producing a protein of the approximate expected size (20kD). The identity of this protein was confirmed by Western blot using the same monoclonal antibodies as in Example 1 (Fig. 4b).

(ii) Construction of Δ E7C/ Δ E6N

Using oligonucleotides (a) in Example 1 and 5'CGCCCGGGTAA-TGTTGTTCCATACTAACTA3' an N-terminal representation of E6 comprising 285bp was amplified from the same HPV-16 clone utilised in Example 1. As well, oligonucleotides 5'CGCCCGGGGAGGAGGAGGATGAAATAGATG3' and

(d) in Example 1 were used to produce a 198bp C-terminal E7 sequence. These were each blunt cloned into pGEM7-Zf(+) (Promega). A fusion cassette was formed by restricting the E6 clone with *KpnI/BgIII* and inserting the E7 sequence upstream as a *KpnI/BamHI* fragment. This fused sequence was then reamplified with *SmaI* and *BgIII* cloning sites for insertion into pGEX-4T-1 containing a termination codon in the GST sequence to produce [GST] $\Delta E7C/\Delta E6Nhh$. After transformation into *E. coli* BL21, protein production was assayed by PAGE followed by Coomassie staining and Western blotting (Fig. 5a and 5b). A protein ($\Delta E7C/\Delta E6Nhh$, represented schematically in Fig.1d) of the expected size (20kD) was evident on Western blots.

EXAMPLE 4

DNA sequencing of E6/E7 full length and deletion constructs

E6/E7 constructs were sequenced in both directions by the dideoxy method using primers that generated overlapping sequence information. The ¹⁷Sequencing™ Kit (Pharmacia) was used to generate ³⁵S-labelled chain-terminated fragments which were analysed on a Sequi-Gen™ (Biorad) electrophoretic gel apparatus. The DNA and corresponding amino acid sequences for E6/E7hh (PPV162.DNA), $\Delta E6C/\Delta E7Nhh$ (CAD600.SEQ) and $\Delta E7C/\Delta E6Nhh$ (C620.TXT) are set out below.

File : PPV162.DNA

Range : 1 - 801 Mode : Normal

Codon Table : Universal

5' ATG CAC CAA AAG AGA ACT GCA ATG TTT CAG GAC CCA CAG GAG CGA CCC AGA AAG

Met His Gln Lys Arg Thr Ala Met Phe Gln Asp Pro Gln Glu Arg Pro Arg Lys

TTA CCA CAG TTA TGC ACA GAG CTG CAA ACA ACT ATA CAT GAT ATA ATA TTA GAA

Leu Pro Gln Leu Cys Thr Glu Leu Gln Thr Thr Ile His Asp Ile Ile Leu Glu

TGT GTG TAC TGC AAG CAA CAG TTA CTG CGA CGT GAG GTA TAT GAC TTT GCT TTT

Cys Val Tyr Cys Lys Gln Gln Leu Leu Arg Arg Glu Val Tyr Asp Phe Ala Phe

CGG GAT TTA TGC ATA GTA TAT AGA GAT GGG AAT CCA TAT GCT GTA TGT GAT AAA

Arg Asp Leu Cys Ile Val Tyr Arg Asp Gly Asn Pro Tyr Ala Val Cys Asp Lys

TGT TTA AAG TTT TAT TCT AAA ATT AGT GAG TAT AGA CAT TAT TGT TAT AGT TTG

Cys Leu Lys Phe Tyr Ser Lys Ile Ser Glu Tyr Arg His Tyr Cys Tyr Ser Leu

TAT GGA ACA ACA TTA GAA CAG CAA TAC AAC AAA CCG TTG TGT GAT TTG TTA ATT

Tyr Gly Thr Thr Leu Glu Gln Gln Tyr Asn Lys Pro Leu Cys Asp Leu Leu Ile

AGG TGT ATT AAC TGT CAA AAG CCA CTG TGT CCT GAA GAA AAG CAA AGA CAT CTG

Arg Cys Ile Asn Cys Gln Lys Pro Leu Cys Pro Glu Glu Lys Gln Arg His Leu

GAC AAA AAG CAA AGA TTC CAT AAT ATA AGG GGT CGG TGG ACC GGT CGA TGT ATG

Asp Lys Lys Gln Arg Phe His Asn Ile Arg Gly Arg Trp Thr Gly Arg Cys Met

- 16 -

TCT TGT TGC AGA TCA TCA AGA ACA CGT AGA GAA ACC CAG CTG CCC GGG ATG CAT

Ser Cys Cys Arg Ser Ser Arg Thr Arg Arg Glu Thr Gln Leu Pro Gly Met His

GGA GAT ACA CCT ACA TTG CAT GAA TAT ATG TTA GAT TTG CAA CCA GAG ACA ACT

Gly Asp Thr Pro Thr Leu His Glu Tyr Met Leu Asp Leu Gln Pro Glu Thr Thr

GAT CTC TAC TGT TAT GAG CAA TTA AAT GAC AGC TCA GAG GAG GAG GAT GAA ATA

Asp Leu Tyr Cys Tyr Glu Gln Leu Asn Asp Ser Ser Glu Glu Glu Asp Glu Ile

GAT GGT CCA GCT GGA CAA GCA GAA CCG GAC AGA GCC CAT TAC AAT ATT GTA ACC

Asp Gly Pro Ala Gly Gln Ala Glu Pro Asp Arg Ala His Tyr Asn Ile Val Thr

TTT TGT TGC AAG TGT GAC TCT ACG CTT CGG TTG TGC GTA CAA AGC ACA CAC GTA

Phe Cys Cys Lys Cys Asp Ser Thr Leu Arg Leu Cys Val Gln Ser Thr His Val

GAC ATT CGT ACT TTG GAA GAC CTG TTA ATG GGC ACA CTA GGA ATT GTG TGC CCC

Asp Ile Arg Thr Leu Glu Asp Leu Leu Met Gly Thr Leu Gly Ile Val Cys Pro

ATC TGT TCT CAG AAA CCA AGA TCT CAT CAC CAT CAC CAT CAC TAA 3'

Ile Cys Ser Gln Lys Pro Arg Ser His His His His His His ***

File : CAD600.SEQ

Range : 1 - 519 Mode : Normal

Codon Table : Universal

54

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Met Gly Asn Pro Tyr Ala Val Cys Asp Lys Cys Leu Lys Phe Tyr Ser Lys Ile
AGT GAG TAT AGA CAT TAT TGT TAT AGT TTG TAT GGA ACA ACA TTA GAA CAG CAA

Ser Glu Tyr Arg His Tyr Cys Tyr Ser Leu Tyr Gly Thr Thr Leu Glu Gln Gln
TAC AAC AAA CCG TTG TGT GAT TTG TTA ATT AGG TGT ATT AAC TGT CAA AAG CCA

I le Asn Cys Gln Lys Pro Tyr Asn Lys Pro Leu Cys Asp Leu Leu Ile Arg Cys
CTG TGT CCT GAA GAA AAG CAA AGA CAT CTG GAC AAA AAG CAA AGA TTC CAT AAT

Leu Cys Pro Glu Glu Lys Gln Arg His Leu Asp Lys Lys Gln Arg Phe His Asn
ATA AGG GGT CGG TGG ACC GGT CGA TGT ATG TCT TGT TGC AGA TCA TCA AGA ACA

Ile Arg Gly Arg Trp Thr Gly Arg Cys Met Ser Cys Cys Arg Ser Ser Arg Thr
CGT AGA GAA ACC CAG CTG CCC GGG ATG CAT GGA GAT ACA CCT ACA TTG CAT GAA

Arg Arg Glu Thr Gln Leu Pro Gly Met His Gly Asp Thr Pro Thr Leu His Glu
TAT ATG TTA GAT TTG CAA CCA GAG ACA ACT GAT CTC TAC TGT TAT GAG CAA TTA

Tyr Met Leu Asp Leu Gln Pro Glu Thr Thr Asp Leu Tyr Cys Tyr Glu Gln Leu
AAT GAC AGC TCA GAG GAG GAG GAT GAA ATA GAT GGT CCA GCT GGA CAA GCA GAA

Asn Asp Ser Ser Glu Glu Glu Asp Glu Ile Asp Gly Pro Ala Gly Gln Ala Glu
CCG GAC AGA GCC CAT TAC AAT ATT GTA ACC TTT TGT TGC AAG TGT GAC TCT ACG

Pro Asp Arg Ala His Tyr Asn Ile Val Thr Phe Cys Cys Lys Cys Asp Ser Thr
CTT CGG AGA TCT CAT CAC CAT CAC CAT CAC TAA 3'

Leu Arg Arg Ser His His His His His His ***

File : C620.TXT

Range : 1 - 519 Mode : Normal

Codon Table : Universal

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Met Glu Glu Asp Glu Ile Asp Gly Pro Ala Gly Gln Ala Glu Pro Asp Arg Ala

CAT TAC AAT ATT GTA ACC TTT TGT TGC AAG TGT GAC TCT ACG CTT CGG TTG TGC

His Tyr Asn Ile Val Thr Phe Cys Cys Lys Cys Asp Ser Thr Leu Arg Leu Cys

GTA CAA AGC ACA CAC GTA GAC ATT CGT ACT TTG GAA GAC CTG TTA ATG GGC ACA

Val Gln Ser Thr His Val Asp Ile Arg Thr Leu Glu Asp Leu Leu Met Gly Thr

CTA GGA ATT GTG TGC CCC ATC TGT TCT CAG AAA CCA GGA TCT CAT ATG CAC CAA

Leu Gly Ile Val Cys Pro Ile Cys Ser Gln Lys Pro Gly Ser His Met His Gln

AAG AGA ACT GCA ATG TTT CAG GAC CCA CAG GAG CGA CCC AGA AAG TTA CCA CAG

Lys Arg Thr Ala Met Phe Gln Asp Pro Gln Glu Arg Pro Arg Lys Leu Pro Gln

TTA TGC ACA GAG CTG CAA ACA ACT ATA CAT GAT ATA ATA TTA GAA TGT GTG TAC

Leu Cys Thr Glu Leu Gln Thr Thr Ile His Asp Ile Ile Leu Glu Cys Val Tyr

TGC AAG CAA CAG TTA CTG CGA CGT GAG GTA TAT GAC TTT GCT TTT CGG GAT TTA

Cys Lys Gln Gln Leu Leu Arg Arg Glu Val Tyr Asp Phe Ala Phe Arg Asp Leu

TGC ATA GTA TAT AGA GAT GGG AAT CCA TAT GCT GTA TGT GAT AAA TGT TTA AAG

Cys Ile Val Tyr Arg Asp Gly Asn Pro Tyr Ala Val Cys Asp Lys Cys Leu Lys

TTT TAT TCT AAA ATT AGT GAG TAT AGA CAT TAT TGT TAT AGT TTG TAT GGA ACA

Phe Tyr Ser Lys Ile Ser Glu Tyr Arg His Tyr Cys Tyr Ser Leu Tyr Gly Thr

ACA TTA AGA TCT CAT CAC CAT CAC CAT CAC TAA 3'

Thr Leu Arg Ser His His His His His His ***

EXAMPLE 5**Immunogenicity of E6/E7hh protein****5 A. Purification of E6/E7hh**

E. coli cells (strain BL21) containing the [GST] E6/E7hh plasmid were induced using 0.1 - 0.5 mM IPTG and harvested 3 - 4 hours after induction. The cells were pelleted by low speed centrifugation and inclusion bodies containing the
10 E6/E7hh protein isolated by sonication and centrifugation. The inclusion pellet was solubilised in 7M Urea or 6M Guanidine HCl and subjected to nickel chelate column chromatography (Porath *et.al.*, *Biochemistry* 22, 1621-1630, 1983). Protein was eluted using either an increasing gradient of imidazole or a decreasing pH
15 gradient, and fractions containing E6/E7hh pooled and dialysed against 25mM Tris, 0.5M NaCl, 1% NOG, 10mM DTT pH7.5. The identity and purity of the dialysed product was determined by Coomassie stained polyacrylamide gel electrophoresis and Western blot using the monoclonal antibodies referred to in Example 1 (Fig. 6a and 6b).

20 B. Immunogenicity of E6/E7hh.

On day 0, two groups of 5 C57BL/6 mice (8 weeks old, female) were inoculated subcutaneously at the base of the tail with 0.1 mL of a formulation containing 6µg ISCOMATRIX™, 19 µg E6/E7hh (purified as in A. above) in PBS pH7.2. A
25 second dose of the formulation was administered at day 14 to group 1, and at day 17 to group 2. At day 21 and 24, mice in groups 1 and 2 (respectively) were bled. Serum antibody responses to E6/E7hh were then measured using the following solid phase EIA:

30 Nunc MaxiSorp EIA plates were coated with E6/E7hh by incubating 0.1 mL/well for 2 hours at 37° of a 10 µg/mL solution in 4M urea in 50mM carbonate buffer, pH 9.5.

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5 The liquid was removed, and the plates were further incubated at 37° for 1 hour with 0.2 mL/well of 1 mg/mL casein in PBS pH 7.2. After 6 washes, 0.1 mL/well of test serum (diluted in PBS pH 7.2, 1 mg/mL casein, 0.5% Tween 20, 0.002% alphazurine A) was added, and the plates incubated for 1 hour at 37°. The plates were then again washed 6x with PBS pH 7.2, 0.5% Tween 20. To detect bound antibody, 0.1mL of 0.1 µg/ml KPL horseradish peroxidase-labelled goat anti-mouse IgG+IgM (H and L chain specific) in PBS pH 7.2, 1 mg/mL casein, 0.5% Tween 20, 0.002% alphazurine A was added to each well. The plates were incubated for 1 hour at 20°, washed 6x with PBS pH 7.2, 0.5% v/v Tween 20, then 0.1 mL of enzyme substrate (3,3',5,5'-tetramethylbenzidine/H₂O₂ formulation, purchased from KPL) was then added. After 10 minutes incubation at 20°, the reaction was stopped by addition of 50 µl of 0.5M H₂SO₄. The coloured product was then measured at 450 nm in a vertical beam spectrophotometer. Titres were expressed as the reciprocal of the serum dilution resulting in an optical density value of 0.1.

25 Table 1 shows that all mice of both groups 1 and 2 produced a significant response following immunization. Titres ranged from 3.17 to 5.66 (expressed in the log₁₀ of the reciprocal dilution resulting in the optical density of 0.1 in the solid phase EIA described above). Pre-existing antibody levels were low or undetectable (measured in sera obtained on day 0 immediately prior to inoculation).

30 Clearly, the E6/E7hh fusion protein is highly immunogenic when administered to mice by this procedure.

As well, E6/E7hh was found to produce specific delayed-type hypersensitivity (DTH) following one dose of a formulation containing E6/E7hh plus ISCOMTM adjuvant. The mice produced specific DTH responses to both E6 and E7 when challenged in the ear with small doses of purified GST-E6 or GST-E7 proteins.

5

Table 1 Log dilution to 0.1 OD.

Group/ mouse	1/1	1/2	1/3	1/4	1/5	2/1	2/2	2/3	2/4	2/5
pre- bleed	<2	<2	<2	<2	<2	<2	2.06	<2	<2	<2
final bleed	3.66	5.66	3.23	3.19	3.79	4.19	4.21	3.71	5.55	3.17

EXAMPLE 6

Transformation studies of E6/E7 gene construct

5

An E6/E7 fusion DNA construct was subcloned into the multiple cloning site of plasmid vector pJ4 Ω (Wilkinson *et al.*, *J. Exp. Med.* (1988) 167:1442-58) as a *Bam*HI fragment to produce pJ4 Ω E6/E7. For comparison purposes pJ4 Ω vectors containing HPV16 E6 (pJ4 Ω E6) and HPV16 E7 (pJ4 Ω E7) ORFs were used. Where neomycin selection was required, the pcDNA3 vector (Invitrogen) containing a neomycin resistance marker was utilised. These plasmids were amplified in *E. coli* and plasmid DNA extracted by alkaline lysis and purified on resin (Qiagen) eluted, ethanol precipitated and resuspended in H₂O. DNA quantity and purity was determined by spectrophotometric measurement at 260 and 280nm. DNA integrity was checked by electrophoresis in 1% agarose gels and ethidium bromide staining. Target cells for transformation were mouse NIH 3T3 cells (CSL Biosciences). The cells were routinely propagated on Minimal Essential Medium (Eagle) supplemented with non-essential amino acids, 2mM glutamine and 10% foetal bovine serum (growth medium).

20

Transfection of NIH 3T3 cells with plasmid DNA was carried out essentially as described in the Promega Technical Bulletin No. 216 using TfxTM-50 to enhance DNA

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uptake. Typical transfection mixtures contained 5µg of test plasmid (pJ4ΩE6/E7, pJ4ΩE7 or pJ4ΩE6 and pJ4ΩE7), and where required 0.1µg pcDNA3. Where pJ4ΩE6 and pJ4ΩE7 were cotransfected 2.5µg of each was used.

- 5 Cells were grown to approximately 80% confluency, the growth medium removed and plasmid DNA mixed with TfxTM-50 in a ratio of 4:1 in Minimal Essential Medium was added and the cells incubated at 37°C.

- Following 1-2 hours incubation at 37°C the transfection mixture was removed and
10 fresh growth medium added. After 48 hours incubation at 37°C transfected cells were removed by trypsinization and either assayed for colony formation in soft agar or incubated for a further 24 hours at 37°C before neomycin selection was applied.

- For assay of colony formation in soft agar the trypsinized cells were resuspended
15 at a density of $1-5 \times 10^5$ cells/mL in RPMI 1640 supplemented with 10% FBS, 2mM glutamine, 10mM Hepes and 0.084% NaHCO₃ (RPMI1640+) and containing 0.4% agarose (Seaplaque low gelling temperature, FMC Bioproducts, USA) maintained at a temperature of 37 °C. Following mixing 2.5mL of this suspension was added to each well of a 6 well tray (Nunc) and allowed to set. The trays were then incubated for a period
20 of 10 - 14 days at 37°C in an atmosphere of 5% CO₂, prior to counting of colonies using an inverted light microscope.

- Selection of neomycin resistant colonies was carried out on subconfluent cell monolayers using RPMI 1640+ containing 700µg/mL neomycin (Geneticin). The
25 monolayers were incubated at 37°C in an atmosphere of 5% CO₂ for 10 - 14 days prior to counting of neomycin resistant colonies using an inverted light microscope. Following counting, the colonies were dispersed by trypsinization and assayed for colony formation in soft agar as described above. The results of a neomycin selection experiment following transfection of 3T3 cells with various plasmid constructs are presented in Table 2.

30

Table 2

Construct (+ pcDNA3)	No. of neomycin resistant colonies	Mean no. of cells per colony
pJ4ΩE6/E7	2	10
pJ4ΩE7	4	>65
pJ4ΩE6 + pJ4ΩE7	11	>66

These results indicate that the E6/E7 fusion is only weakly transforming in comparison with E7 or E6 + E7. Both colony numbers and cell growth for the E6/E7 fusion were low in comparison with the unfused wild-type sequences. This indicates
5 that the outcome of fusing the E6 and E7 sequences is impairment of the ability of these sequences to promote cell transformation.

Persons skilled in this art will appreciate that variations and modifications may be made to the invention as broadly described herein, other than those specifically
10 described without departing from the spirit and scope of the invention. It is to be understood that this invention extends to include all such variations and modifications.

CLAIMS:

1. A variant of the human papillomavirus (HPV) E6 or E7 protein, said variant being able to elicit a humoral and/or cellular immune response against HPV in a host animal but not being cell-transforming in said host animal.
2. A variant according to claim 1, which is a deletion mutant of the HPV E6 or E7 protein comprising at least 50% of the full-length sequence of the E6 or E7 protein.
3. A variant according to claim 2, wherein said deletion mutant corresponds to approximately two-thirds of full length sequence of the E6 or E7 protein.
4. A variant according to claim 3 wherein said deletion mutant corresponds to the N-terminal two-thirds of the full length sequence of the E6 or E7 protein.
5. A variant according to claim 3 wherein said deletion mutant corresponds to the C-terminal two-thirds of the full length sequence of the E6 or E7 protein.
6. A variant according to claim 1, which is a fusion protein comprising E6 and/or E7 protein moieties, and optionally a linkage moiety.
7. A variant according to claim 6, wherein said fusion protein comprises an E6 and/or E7 protein moiety fused or otherwise coupled to a foreign protein or peptide moiety.
8. A variant according to claim 7, wherein said foreign protein or peptide moiety is selected from protein or peptide moieties to assist in purification of the fusion protein or protein or peptide moieties to enhance the immunogenicity of the fusion protein.

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9. A variant according to any of claims 6 to 8 wherein the E6 and/or E7 protein moieties are selected from full length E6 or E7 protein moieties or non-full length deletion mutants thereof.
10. A variant according to any of claims 6 to 9 wherein said linkage moiety comprises from 1 to 50, preferably from 1 to 20, and most preferably from 1 to 5 amino acid residues.
11. A variant according to claim 6, which comprises a fusion protein comprising a full length E6 protein moiety fused to a full length E7 protein moiety.
12. A variant according to claim 6, which comprises a fusion protein comprising a first protein moiety which is the N-terminal two-thirds of the E6 protein fused to a second protein moiety which is the C-terminal two-thirds of the E7 protein.
13. A variant according to claim 6, which comprises a fusion protein comprising a fusion protein comprising a first protein moiety which is the N-terminal two-thirds of the E7 protein fused to a second protein moiety which is the C-terminal two-thirds of the E6 protein.
14. A variant according to any of claims 1 to 13, wherein said E6 or E7 protein is selected from the HPV-16, HPV-18, HPV-6 and HPV-11 genotypes.
15. A variant according to claim 14, wherein said E6 or E7 protein is selected from the HPV-16 and HPV-18 genotypes.
16. A vaccine composition for use in eliciting a humoral and/or cellular immune response against HPV in a host animal, said composition comprising a variant of the HPV E6 or E7 protein according to any of claims 1 to 15, together with a pharmaceutically acceptable carrier and/or diluent.
17. A vaccine composition according to claim 16, further comprising an adjuvant.

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18. A method for eliciting a humoral and/or cellular response against HPV in a host animal, which method comprises administering to the host animal an effective amount of a variant of the HPV E6 or E7 protein according to any of claims 1 to 15.
19. A method according to claim 18, wherein said variant of the HPV E6 or E7 protein is administered in a composition together with a pharmaceutically acceptable carrier and/or diluent.
20. A method according to claim 19, wherein said composition further comprises an adjuvant.
21. A method according to any of claims 18 to 20, wherein said host animal is a human.
22. Use of a variant of the HPV E6 or E7 protein according to any of claims 1 to 15, in eliciting an immune response against HPV in a host animal.

FIG 1(a) GST E6 E7 hh



FIG 1(b) E6 E7 hh



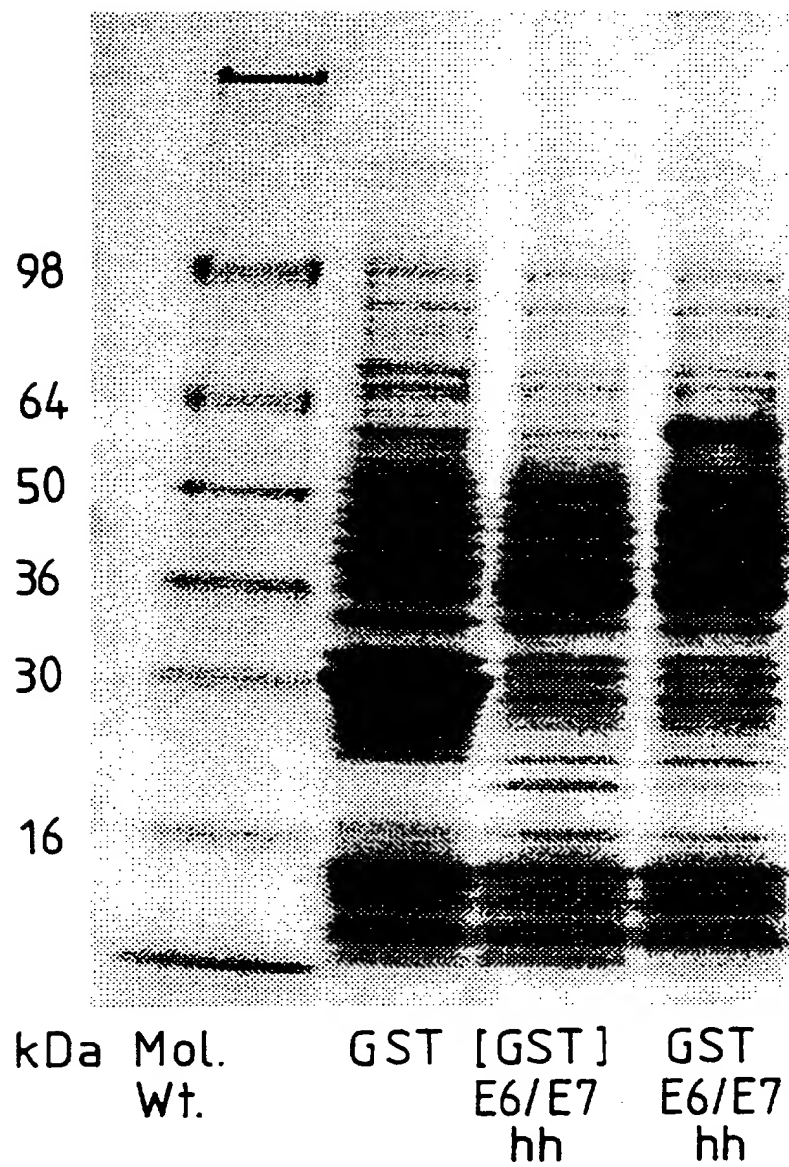
FIG 1(c) Δ E6C Δ E7N hh



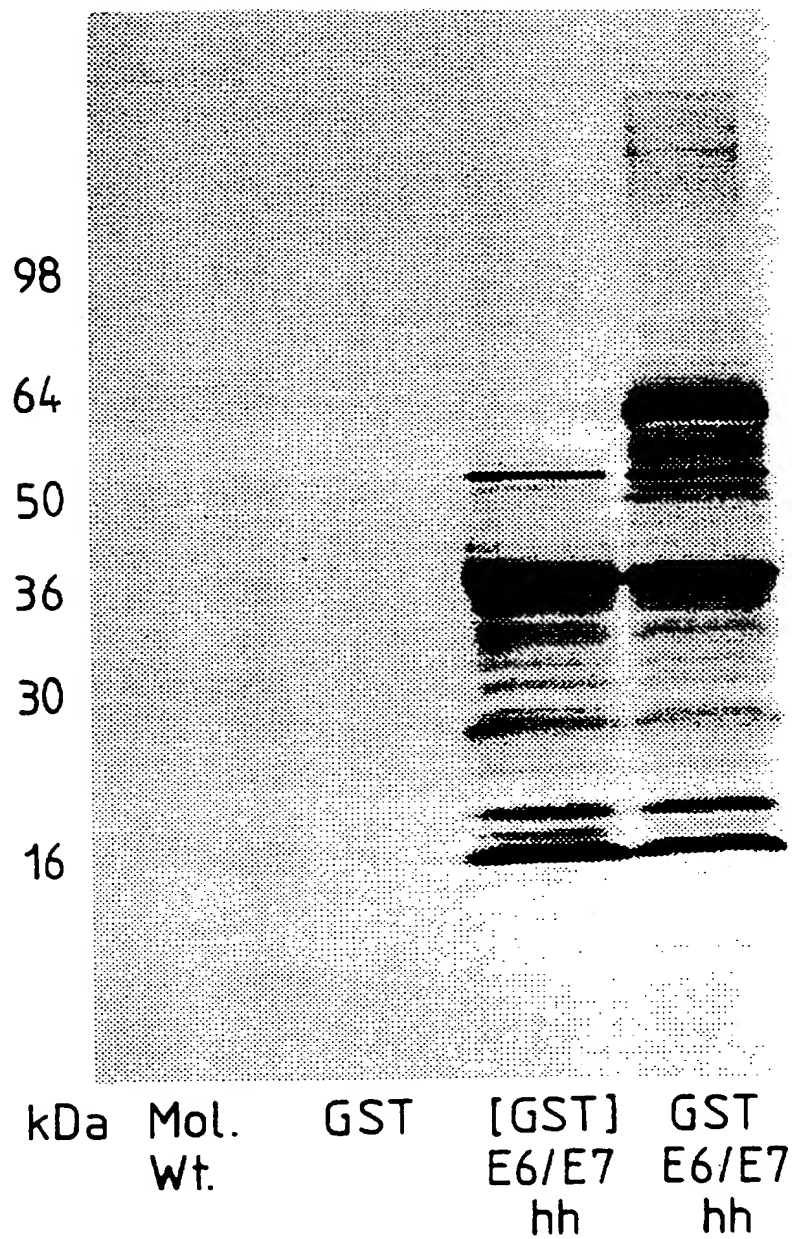
FIG 1(d) Δ E7C Δ E6N hh



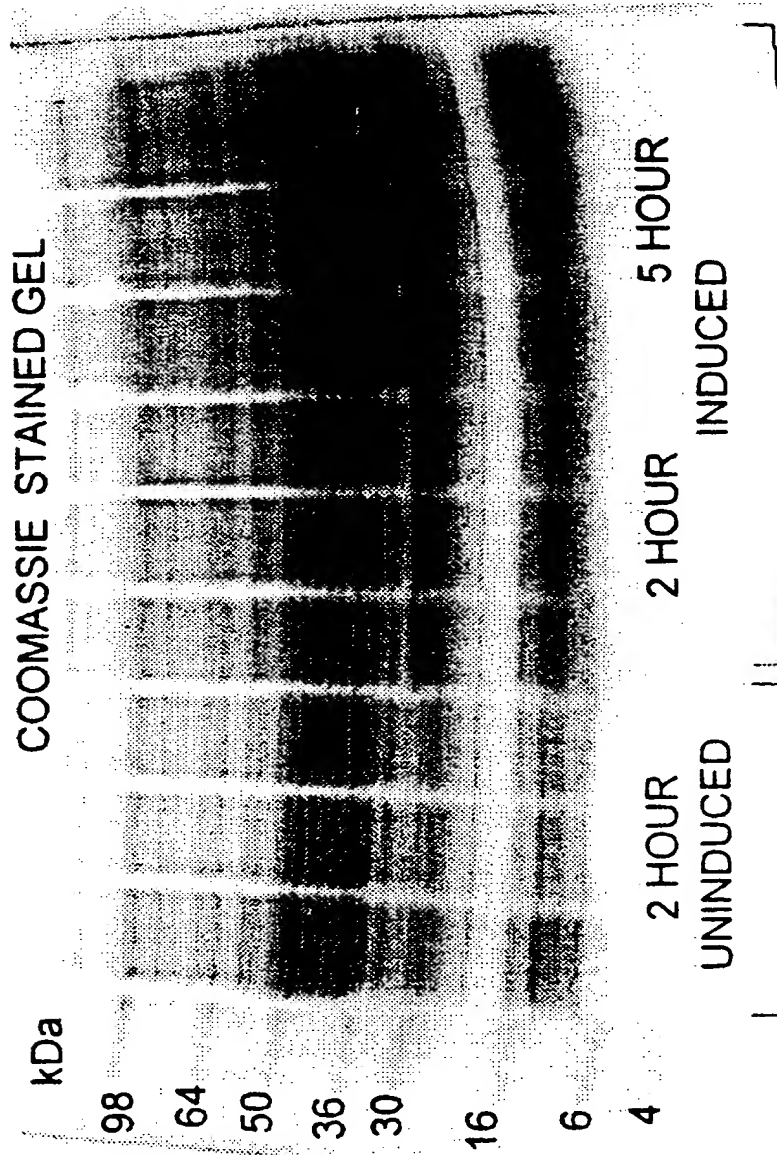
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FIG 2

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FIG 3

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FIG 4a

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WESTERN BLOT



FIG 4b

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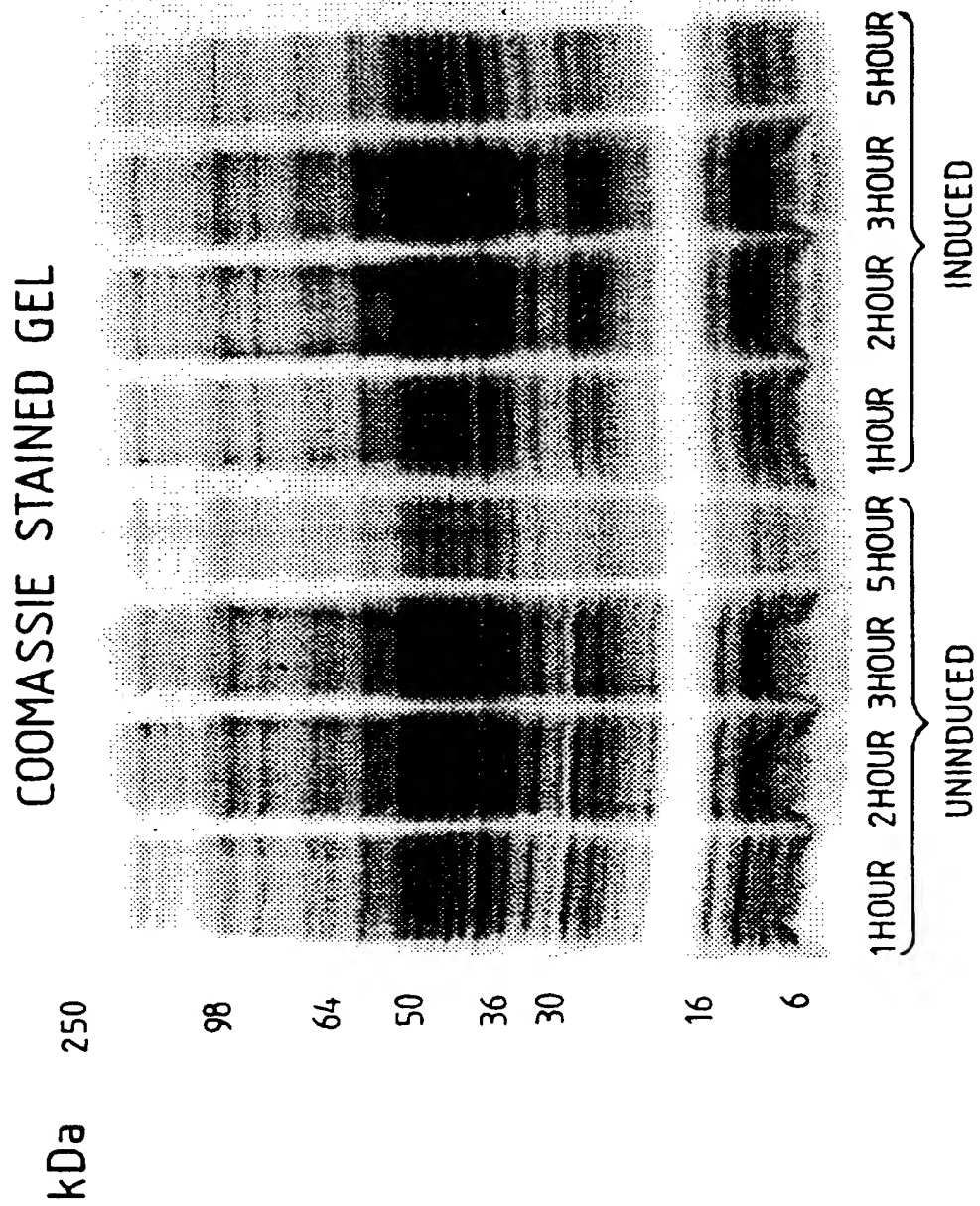


FIG 5a

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FIG 5b

WESTERN BLOT

kDa

250

98

64

50

36

30

16

6

4

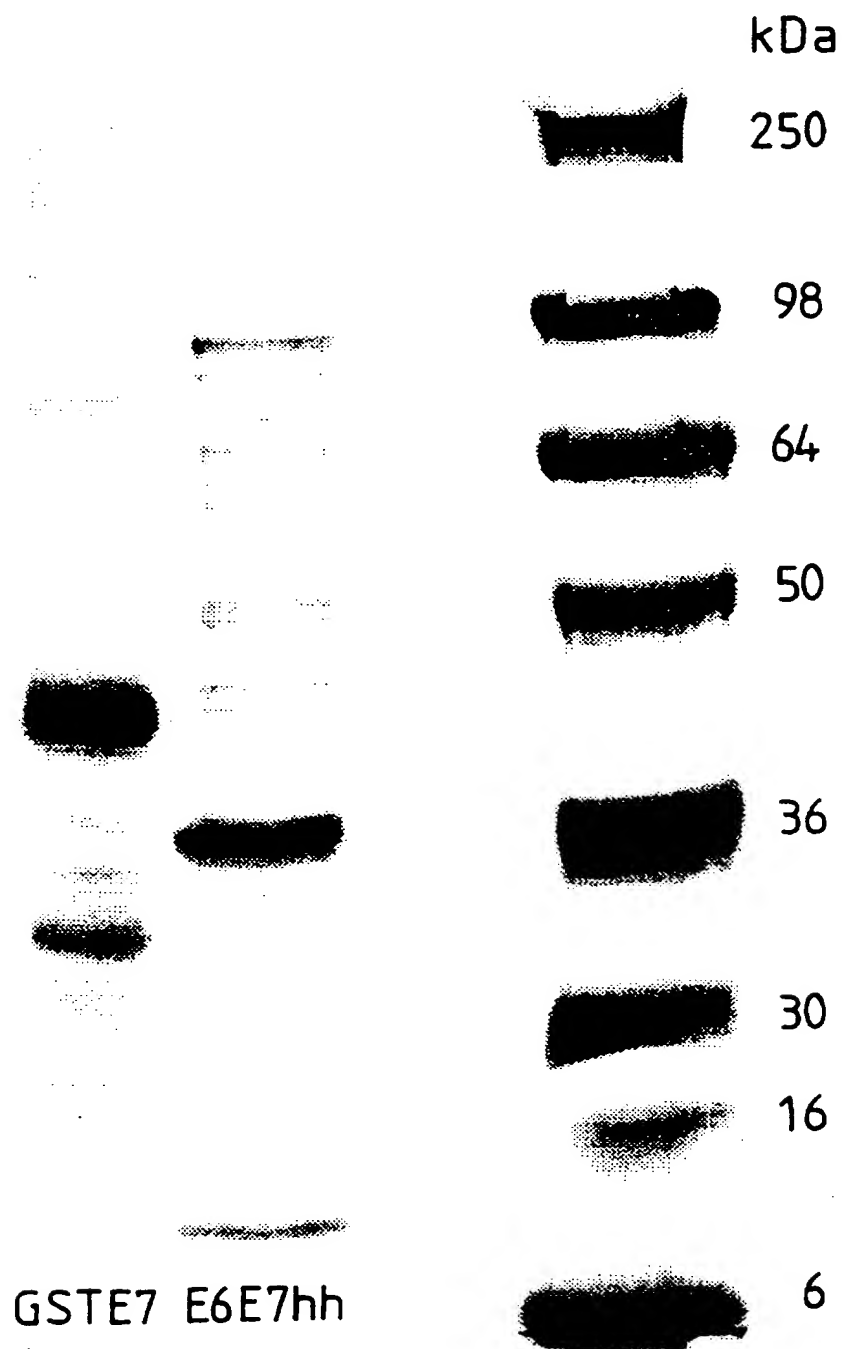
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	1 HOUR		3 HOUR	
INDUCED		INDUCED		INDUCED
1 HOUR		2 HOUR		4 HOUR

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FIG 6a

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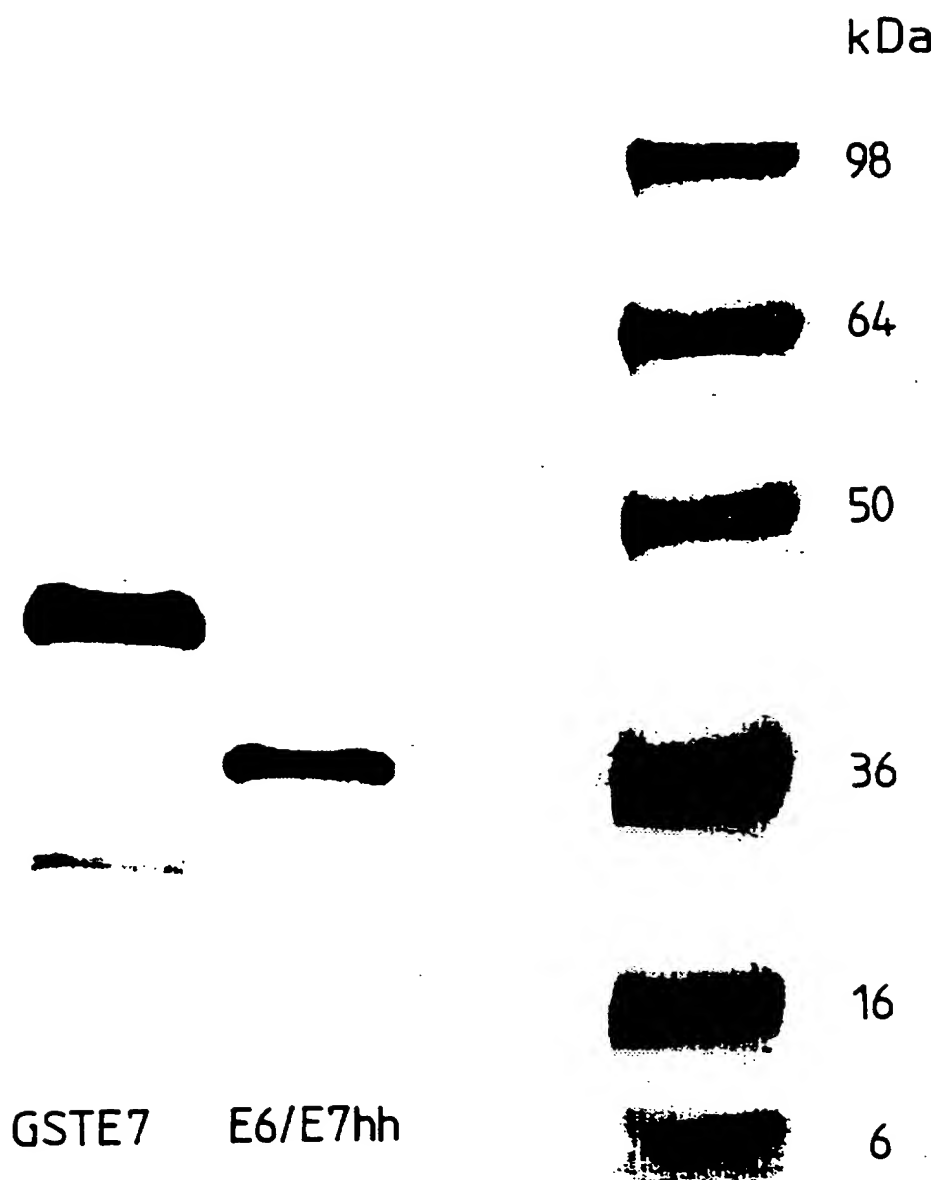


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FIG 6b

Western Blot Probed with MAb anti E7



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A. CLASSIFICATION OF SUBJECT MATTER				
Int Cl ⁶ : C07K 14/025; C12N 15/37; C12N 15/62; A61K 39/12				
According to International Patent Classification (IPC) or to both national classification and IPC				
B. FIELDS SEARCHED				
Minimum documentation searched (classification system followed by classification symbols) WPAT : Chemical Abstracts: keywords as below				
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched				
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) WPAT : JAPIO : Keywords : human papilloma virus or HPV and E6 or E7 CAS online : Keywords as above and indexing terms antigen? or immun? and 1991-1996				
C. DOCUMENTS CONSIDERED TO BE RELEVANT				
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.		
X	WO,A, 92/10513 (THE UNIVERSITY OF QUEENSLAND AND CSL LIMITED) 25 June 1992. See entire document	1, 6-10, 14-22		
X	WO,A, 92/05248 (BRISTOL-MYERS SQUIBB COMPANY) 2 April 1992. See entire document	1, 6-10, 14-22		
X	AU,B, 76212/91 (650648) (BEHRINGWERKE AKTIENGESELLSCHAFT) 14 November 1991. See entire document	1, 6, 7, 8-10, 14-22		
<input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C <input checked="" type="checkbox"/> See patent family annex				
<table border="0" style="width: 100%;"> <tr> <td style="vertical-align: top; width: 50%;"> <p>* Special categories of cited documents:</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </td> <td style="vertical-align: top; width: 50%;"> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</p> <p>"&" document member of the same patent family</p> </td> </tr> </table>			<p>* Special categories of cited documents:</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p>	<p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</p> <p>"&" document member of the same patent family</p>
<p>* Special categories of cited documents:</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p>	<p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</p> <p>"&" document member of the same patent family</p>			
Date of the actual completion of the international search 3 April 1996		Date of mailing of the international search report 15TH APRIL 1996.		
Name and mailing address of the ISA/AU AUSTRALIAN INDUSTRIAL PROPERTY ORGANISATION PO BOX 200 WODEN ACT 2606 AUSTRALIA Facsimile No.: (06) 285 3929		Authorized officer KAREN AYERS Telephone No.: (06) 283 2082		

INTERNATIONAL SEARCH REPORT

International Application No.

PCT/AU 95/00868

C (Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	Intervirolgy, Vol, 37, 1994, H. Stoppler <u>et al.</u> , "Transforming proteins of the papillomaviruses" pages 168-179, see entire document.	1-22

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INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No.
PCT/AU 95/00868

This Annex lists the known "A" publication level patent family members relating to the patent documents cited in the above-mentioned international search report. The Australian Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent Document Cited in Search Report				Patent Family Member			
WO,A	92/10513	AU EP	90709/91 561885	AU JP	660954 6503559	CA	2097916
WO,A	92/05248	AU	87629/91	CN	1067382		
AU,B	76212/91, (650648)	EP JP	456197 4227000	CA PT	2042236 97621	DE	4015044